

AD _____

Award Number: **DAMD17-96-1-6229**

TITLE: **Cancers Missed on Mammography**

PRINCIPAL INVESTIGATOR: **Robert M. Nishikawa, Ph.D.**

CONTRACTING ORGANIZATION: **University of Chicago**
Chicago, Illinois 60637

REPORT DATE: **October 2000**

TYPE OF REPORT: **Annual**

PREPARED FOR: **U.S. Army Medical Research and Materiel Command**
Fort Detrick, Maryland 21702-5012

DISTRIBUTION STATEMENT: **Approved for public release; Distribution**
Unlimited

The views, opinions and/or findings contained in this report are those of the author(s) and should not be construed as an official Department of the Army position, policy or decision unless so designated by other documentation.

DTIC QUALITY INSPECTED 4
20010124 018

REPORT DOCUMENTATION PAGEForm Approved
OMB No. 074-0188

Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing this collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204, Arlington, VA 22202-4302, and to the Office of Management and Budget, Paperwork Reduction Project (0704-0188), Washington, DC 20503

1. AGENCY USE ONLY
(Leave blank)**2. REPORT DATE**
October 2000**3. REPORT TYPE AND DATES COVERED**
Annual (23 Sep 99 - 22 Sep 00)**4. TITLE AND SUBTITLE**
Cancers Missed on Mammography**5. FUNDING NUMBERS**
DAMD17-96-1-6229**6. AUTHOR(S)**
Robert M. Nishikawa, Ph.D.**7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES)**
University of Chicago
Chicago, Illinois 60637**E-MAIL:**
r-nishikawa@uchicago.edu**8. PERFORMING ORGANIZATION**
REPORT NUMBER**9. SPONSORING / MONITORING AGENCY NAME(S) AND ADDRESS(ES)**
U.S. Army Medical Research and Materiel Command
Fort Detrick, Maryland 21702-5012**10. SPONSORING / MONITORING**
AGENCY REPORT NUMBER**11. SUPPLEMENTARY NOTES****12a. DISTRIBUTION / AVAILABILITY STATEMENT**
Approved for public release; distribution unlimited**12b. DISTRIBUTION**
CODE**13. ABSTRACT (Maximum 200 Words)**

Observer error in reading screening mammograms has been identified as a significant factor in delayed diagnosis of breast cancer. The magnitude of the problem is estimated to be about 30% of potentially detectable cancers are overlooked for one or more years before detection. Computer-aided diagnosis (CAD) programs have been developed to aid radiologists in the detection task, and pre-clinical studies have shown that CAD applied to digitized mammography films can flag about 50% of radiologists' observational oversights. Preliminary study has also shown a wide variability in radiologist observer performance. The purpose of this investigation is to test how many additional cancers are detected by radiologists using CAD, in an observer study using an enriched mixture of cancers. A pilot study has been completed and the results were used to plan the full observer study: 400 cases containing 75 cancers with 6 radiologists. Final case selection is being done and observers are being recruited.

14. SUBJECT TERMS
Mammography, computer-aided diagnosis, missed cancers**15. NUMBER OF PAGES**
11**16. PRICE CODE****17. SECURITY**
CLASSIFICATION
OF REPORT
Unclassified**18. SECURITY**
CLASSIFICATION
OF THIS PAGE
Unclassified**19. SECURITY**
CLASSIFICATION
OF ABSTRACT
Unclassified**20. LIMITATION OF**
ABSTRACT
Unlimited

Table of Contents

| | |
|------------------------------------|---|
| Cover..... | 1 |
| SF 298..... | 2 |
| Table of Contents | 3 |
| Introduction | 4 |
| Body | 4 |
| Key Research Accomplishments | 6 |
| Reportable Outcomes | 7 |
| Conclusions | 7 |
| References..... | 7 |
| Appendices | 9 |

4. INTRODUCTION

Double reading of mammograms has been shown to significantly increase the number of cancers detected.¹⁻⁵ Computer-aided diagnosis (CAD) has been proposed as an efficient method of implementing double reading.⁶ For CAD to be effective computers must find cancers that are missed by radiologists, and radiologists must react appropriately to the computer prompts. Others and we have found that computer detection schemes can find over 50% of the observational misses made by radiologists reading mammograms.⁷⁻⁹ Our current study is designed to show that CAD can help detect cancers that they might otherwise be overlooked. We will collect a large database of cancers already missed by radiologists in routine clinical practice, and will test observers without and with the aid of CAD. It is expected that radiologists will detect about 10 to 15% more cancers using CAD, which would have important implications for bringing this technique into clinical practice. We will also learn much more about the reasons for and types of radiologist misses on mammography.

5. BODY OF REPORT

5.1 Tasks

There are five tasks in the Statement of Work, which are listed below.

Task 1. Preparation of review forms and finalization of eligibility characteristics for cases to be entered into the missed lesion database.

Task 2. Accumulation of database cases and copying/digitizing 100 missed malignant cases and 300 normal cases, with categorization of features and characteristics of the malignant case. Verification of missed lesion cases. Ongoing data entry.

Task 3. Computer runs producing hard copy of computer output for use in observer experiment and preparation of cases for observer experiment. Ongoing data entry of computer accuracy and truth table for missed lesion database. Final design of details of observer performance study.

Task 4. An observer experiment conducted on 15 observers at about 3 hours per session, with 6 sessions per observer spaced at 2-3 months apart. Goal is to perform 2 observation sessions and analysis minimum per week, entering observation data into a computer database. Ongoing data entry.

Task 5. Final analysis of data comparing CAD observer results with non-CAD results and observer variability, and preparation of report summarizing the results of the observer experiment and the clinical characteristics of the missed lesions.

5.1.1 Preparation of forms

A copy of the review form has been submitted previously. The eligibility criteria are as follows:

1. Patients who have had screen-film mammograms read at the participating mammography facilities.
2. For cases of missed lesions, the mammogram had to be read clinically as normal in the area where a cancer subsequently developed, and the error had to be one of observation (failure to see the lesion) rather than interpretation (seeing the lesion and categorizing it as benign). In cases where the cancer is visible on multiple examinations prior to diagnosis, the two expert mammographers reviewing the cases will collaboratively select a single representative screening exam as the index missed case.
3. Case is a minimum of 1 year old (to avoid any interference with clinical care), unless bilateral mastectomy has been performed, or unless films clinically equivalent to those entered into the study from other years are available.
4. Case is not involved in any medical-legal action.
5. No copy films will be used that include significant marks made by a previous observer prior to the copying, and no originals with such permanent marks will be used.

5.1.2 Development of database of missed lesions

The database is nearly complete. All 100 cases with a missed cancer have been identified, although not all have been digitized or categorized. Over half the normals have been collected, with 160 cases from the University of Chicago. The remaining normals will be collected from the University of New Mexico and the University of Chicago. Three tables in the Appendices summarize some of the characteristics of the cancers entered into our database. The average size of the cancers is 11.7 mm.

5.1.3 Computer analysis of case

We will run the computer CAD program on the database, once the database has been completed. This will allow us to use the most current version of our detection schemes. It will take approximately 1 week to run and print the computer results.

5.1.4 Observer study

The formal observer study has not yet begun. We have just complete a pilot observer study using 75 cases that contained 24 cancers (all but two were clinically missed cancers) and 51 normals. The objective was to gather data as to the minimum number of cancers we need to include in our observer study and to examine if the computer false positive rate was going to be too high. Also, we tested the logistics of the planned observer study: the user interface to record observers' ratings, whether the questions asked were understandable by the radiologists and how effective was our training session.

Five radiologists read the cases in one session. For each case, they answered two questions: (i) Give your BI-RADS assessment of this case; and (ii) what is your level of confidence that the patient should be called back for further work-up or a biopsy? The later question was answered using a visual analog scale, in which the observer marks a point on a 5-cm line – the left end of the line is labeled “definitely DO NOT call back”, and the right end is marked “Definitely call back”. These questions were first answered after the radiologists viewed

the films and a second time after viewing the computer detection output. Some minor “bugs” in the software have been identified and will be corrected. Otherwise the interface was easy to use and recorded all the information that we needed.

The fifth observer has not yet finished the study. The results for the other four are given in Table V. The important information from this experiment in terms of planning the full observer study are: A_Z with and without aid and the correlation between the two A_Z values. It was disappointing that we did not see much of an improvement when the readers used the computer aid. We attribute this to:

1. High false-positive rate of the computer aid (approximately 2.5 per image). The sensitivity for this set of images was roughly 55%, compared to the clinical reading of 8%. The high false-positive rate reduces the time the radiologist spends considering the computer findings and therefore, reduces the likelihood that an overlooked cancer detected by the computer will be noticed. Furthermore, the high false-positive rate increased radiologists’ call back rate, thus reducing performance. To solve this, we will further optimize our algorithms. The computer results we used in this study were from our clinical study. The clinical study used algorithms developed in 1994.

2. Insufficient training. Two of the readers had used the R2 Imagechecker CAD system. This biased them to pay more attention to the calcification results and spend less time considering the mass results. Since most of missed cancers were masses, the radiologists were biased against finding the missed masses. A more extensive and interactive training regime will be developed.

Given that we expect to measure an improvement when the radiologists use CAD, we will power our experiment to measure a 0.06 difference in A_Z . Table VI shows the power for $\alpha=0.05$, given the total number of cases, the number of cancer cases, the without aid A_Z , and the correlation between the two A_Z values. The power calculation is most sensitive to the correlation between A_Z values, followed by the number of cancer cases. These calculations are assuming a single observer. When multiple observers are used, the power goes up depending on how similar, in terms of A_Z , the readers are.

One of the objectives of our proposal is to simulate, as best as possible, actual reading conditions. To do this we would like to use a low cancer prevalence in our observer study. This is an attempt to require the readers to maintain high vigilance in reading as need to do clinically where the call back rate is 5-15%. We will use 75 cancers and 400 cases which gives a prevalence of 19%. We will have six readers, which should give use sufficient power to see at least a 0.06 difference in A_Z . If we are not able to measure a statistically significant difference with six readers, we plan to use up to six additional readers.

5.1.5 Data Analysis

Data analysis of the missed lesion/CAD study cannot begin until the observer study has been completed.

5.2 Discussion

Given the results from our pilot study, we have changed our observer study to include 400 cases that contain 75 cancers and 6 observers. This should be sufficient to see an improvement in A_z of 0.06 when CAD is used. We are now finalizing case selection. We will begin recruiting observers and start the observer study.

5.3 Recommendations in relation to the Statement of Work

- Other than re-specifying the number of cancer cases and the number of observers we will use in our observer study, we do not anticipate making any changes to the Statement of Work.

6. KEY RESEARCH ACCOMPLISHMENTS

- Pilot observer study performed
- Detailed planning of observer study complete
- Final case selection and observer recruitment are being made.

7. REPORTABLE OUTCOMES

None since last report.

8. CONCLUSIONS

Data collection is nearly complete and so we will begin to conduct our main observer study in year 2001. Valuable data has been collected from a preliminary smaller observer study, which will influence the design of the larger scale observer study. We anticipate that we will be able to demonstrate that CAD can reduce the number of missed cancers by 50%, which has not yet been shown in a structured observer experiment. These results should provide information on which health care providers and governmental organizations can base decisions on the value of introducing this promising new technology into the clinical practice of breast cancer screening.

9. REFERENCES

1. C. E. Metz and J. H. Shen, "Gains in accuracy from replicated readings of diagnostic images: prediction and assessment in terms of ROC analysis," *Medical Decision Making* **12**, 60-75 (1992).
2. E. L. Thurffjell, K. A. Lernevall and A. A. S. Taube, "Benefit of independent double reading in a population-based mammography screening program," *Radiology* **191**, 241-244 (1994).
3. R. M. L. Warren and S. W. Duffy, "Comparison of single reading with double reading of

- mammograms, and change in effectiveness with experience," *British Journal of Radiology* **68**, 958-962 (1995).
4. C. A. Beam, D. C. Sullivan and P. M. Layde, "Effect of human variability on independent double reading in screening mammography," *Academic Radiology* **3**, 891-897 (1996).
 5. J. Brown, S. Bryan and R. Warren, "Mammography screening: An incremental cost effectiveness analysis of double versus single reading of mammograms," *British Medical Journal* **312**, 809-812 (1996).
 6. R. A. Schmidt and R. M. Nishikawa, "Digital screening mammography," *Principles & Practice of Oncology* **8**, 1-16 (1994).
 7. R. A. Schmidt, R. M. Nishikawa, R. Osnis, K. L. Schreibman, M. L. Giger and K. Doi, "Computerized detection of lesions missed by mammography," in *Digital Mammography '96*, edited by K. Doi, M. L. Giger, R. M. Nishikawa and R. A. Schmidt (Elsevier Science, Amsterdam, 1996), pp. 105-110.
 8. J. Roehrig, T. Doi, A. Hasegawa, B. Hunt, J. Marshall, H. Romsdahl, A. Schneider, R. Sharbaugh and W. Zhang, "Clinical results with R2 Imagechecker system," in *Digital Mammography - Nijmegen, 1998*, edited by N. Karssemeijer, M. Thijssen, J. Hendriks and L. van Erning (Kluwer Academic Publishers, Dordrecht, The Netherlands, 1998), pp. 395-400.
 9. G. M. te Brake, N. Karssemeijer and J. H. C. L. Hendriks, "Automated detection of breast carcinomas not detected in a screening program," *Radiology* **207**, 465-471 (1998).
 10. R.A. Schmidt, G.M. Newstead, M.N. Linver, G.W. Eklund, C.E. Metz, M.A. Winkler, R.M. Nishikawa, "Mammographic Screening: Sensitivity of General Radiologists," in *Digital Mammography - Nijmegen, 1998*, edited by N. Karssemeijer, M. Thijssen, J. Hendriks and L. van Erning (Kluwer Academic Publishers, Dordrecht, The Netherlands, 1998), pp. 383-388.

10. APPENDICES

Table 1. Distribution of breast density in our database

| Breast Density | Frequency of Occurrence |
|----------------|-------------------------|
| Normal | 0.30 |
| Fatty | 0.21 |
| Dense | 0.37 |
| Focal | 0.09 |

Table 2. Distribution of subtlety on a 5-point scale, where 1 is extremely subtle.

| Subtlety Rating | Frequency of Occurrence |
|-----------------|-------------------------|
| 1 | 0.16 |
| 2 | 0.39 |
| 3 | 0.37 |
| 4 | 0.05 |
| 5 | 0 |

Table 3. Distribution by lesion type*

| Type of Lesion | Frequency of Occurrence |
|------------------------|-------------------------|
| Asymmetric Density | 0.29 |
| Archectural Distortion | 0.24 |
| Developing Density | 0.07 |
| Mass | 0.46 |
| Calcifications | 0.10 |

*numbers sum to greater than 1, because some cases have multiple lesions.

Table 4. Distribution of possible reasons for cancers being missed.*

| Possible Reason | Frequency of Occurrence |
|------------------------------|-------------------------|
| Seen on only 1 view | 0.48 |
| Obscured by overlying tissue | 0.40 |
| Looks like normal tissue | 0.36 |
| "Busy" breast | 0.29 |
| Film technique | 0.26 |
| Distracting lesions | 0.24 |
| Subtle lesion | 0.14 |
| Marginal lesion | 0.10 |
| Developing density | 0.10 |
| Benign appearing lesion | 0.07 |
| Lack of prior films | 0.07 |
| Too small to prompt workup | 0.05 |
| Lucent lines | 0.05 |
| Stable lesion | 0.02 |

*numbers sum to greater than 1, because up to three reasons were given per case.

Table V. Summary from pilot observer study.

| Reader | Unaided | With Aid | Correlation between aid and no aid |
|--------|---------|----------|------------------------------------|
| A | 0.686 | 0.685 | 0.967 |
| B | 0.725 | 0.775 | 0.817 |
| C | 0.805 | 0.793 | 0.943 |
| D | 0.710 | 0.688 | 0.988 |
| mean | 0.731 | 0.735 | 0.929 |

Table VI. The power, for $\alpha=0.05$, to show a difference in A_z of 0.06 under different conditions.

| Power | Total Number of Cases | Number of Cancer Cases | A_z of Unaided Reading | Correlation between aid and no aid |
|-------|-----------------------|------------------------|--------------------------|------------------------------------|
| 0.85 | 400 | 100 | 0.73 | 0.82 |
| 0.80 | 400 | 84 | 0.73 | 0.82 |
| 0.96 | 400 | 84 | 0.73 | 0.90 |
| 0.80 | 400 | 46 | 0.73 | 0.90 |
| 0.78 | 400 | 46 | 0.70 | 0.90 |
| 0.86 | 400 | 46 | 0.80 | 0.90 |